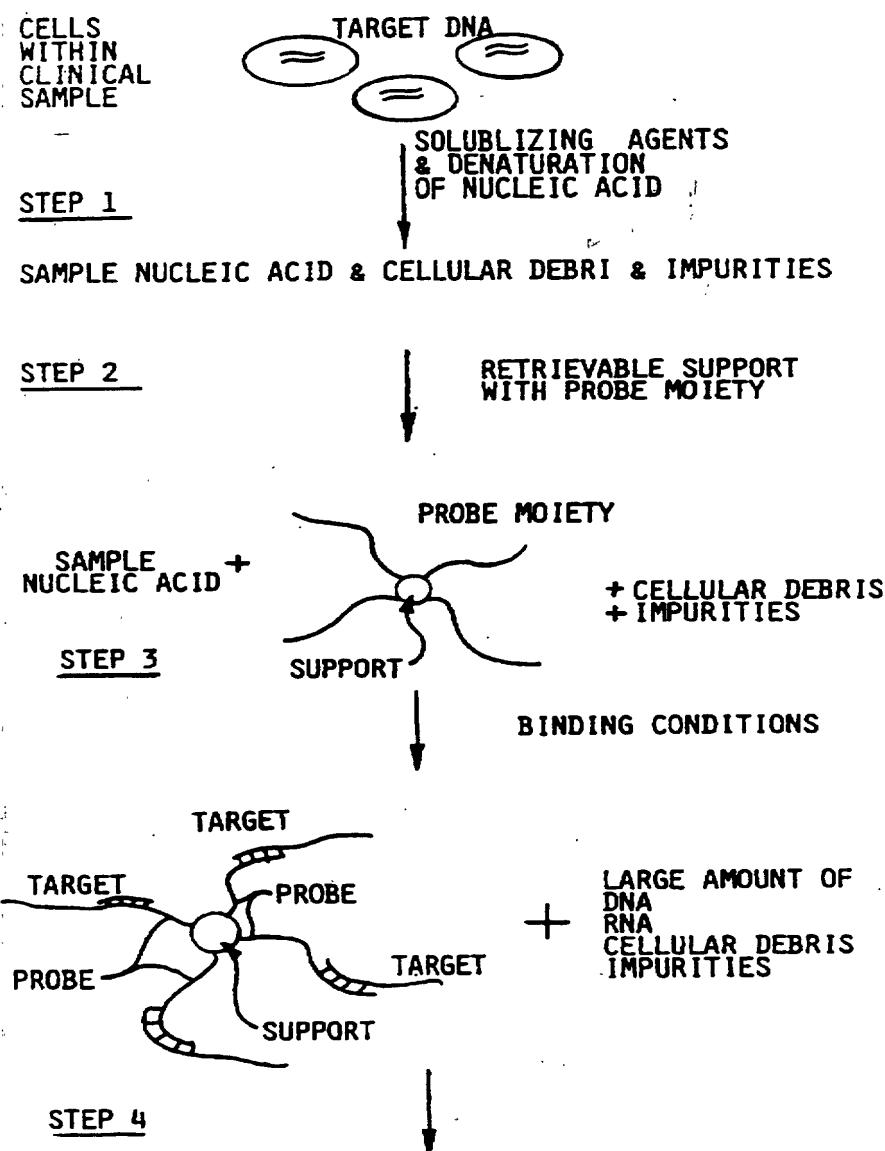


FIG. 1a



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FIG. I b

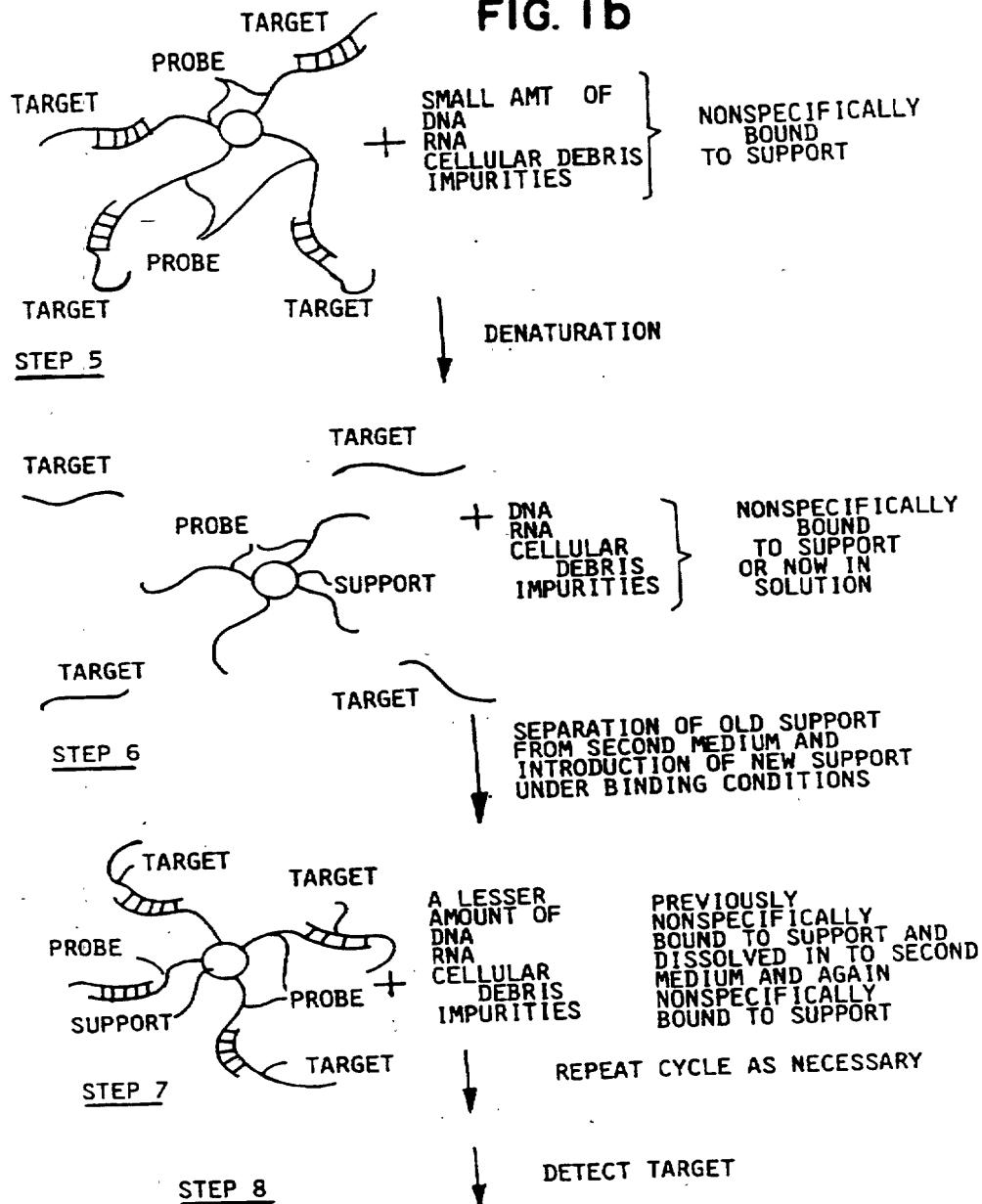


FIG. 2a

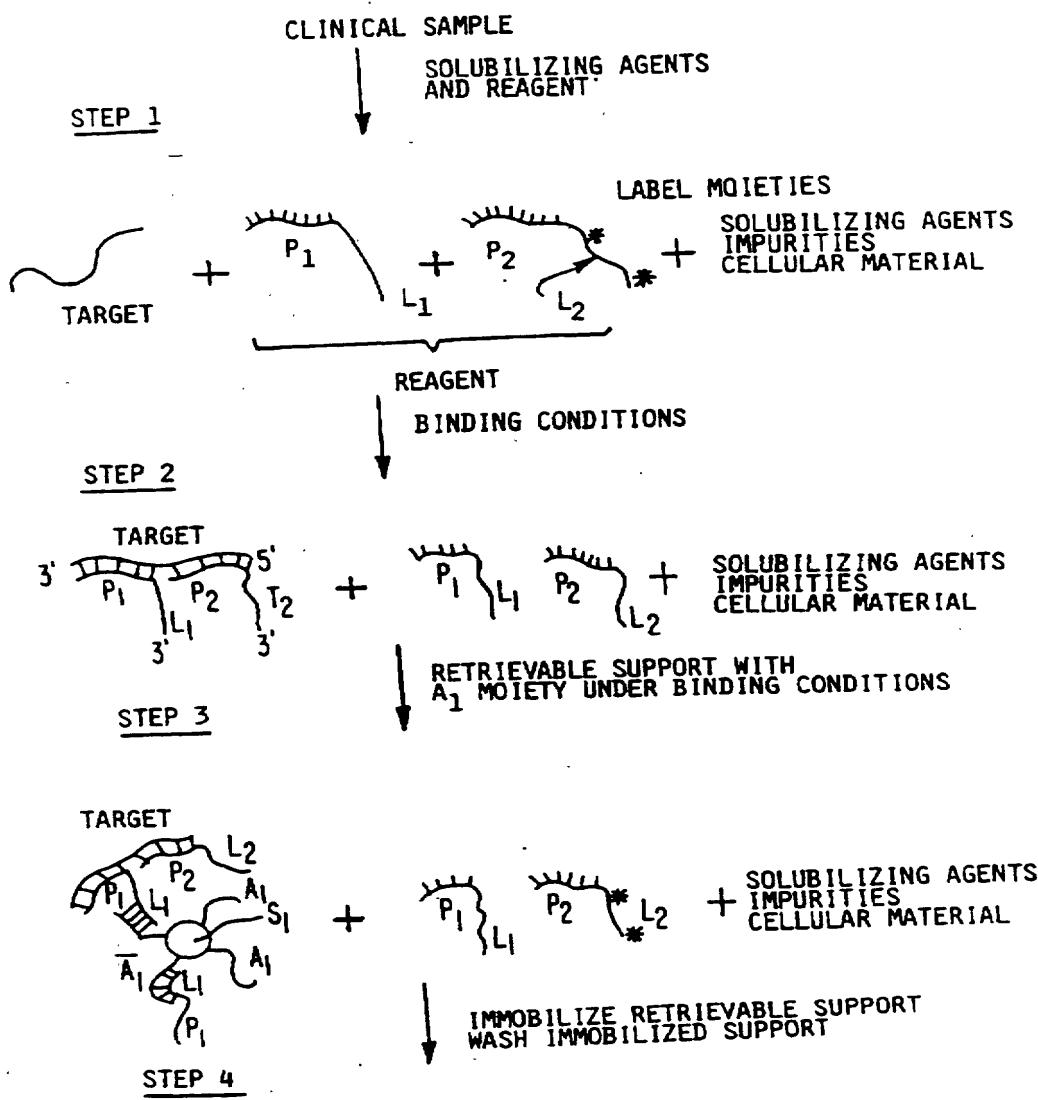
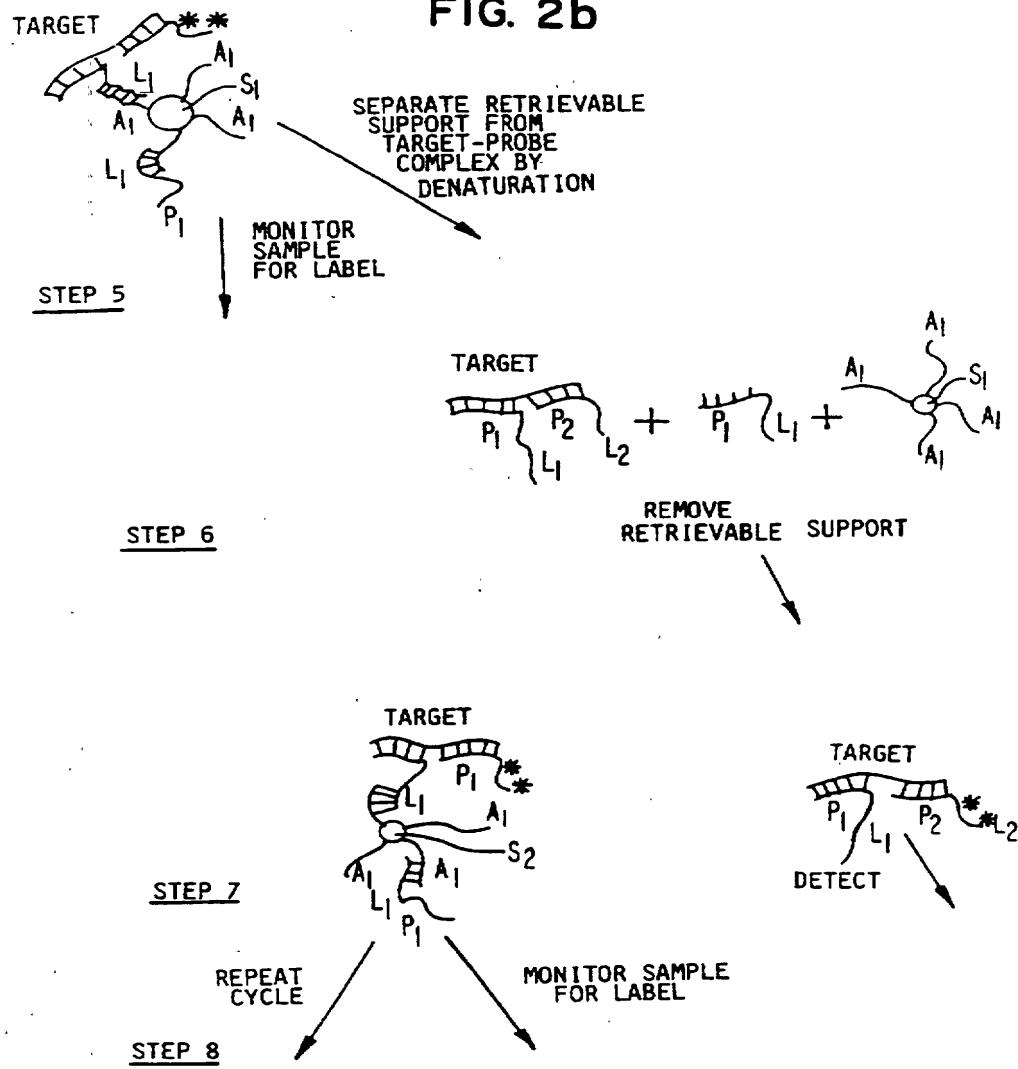


FIG. 2b



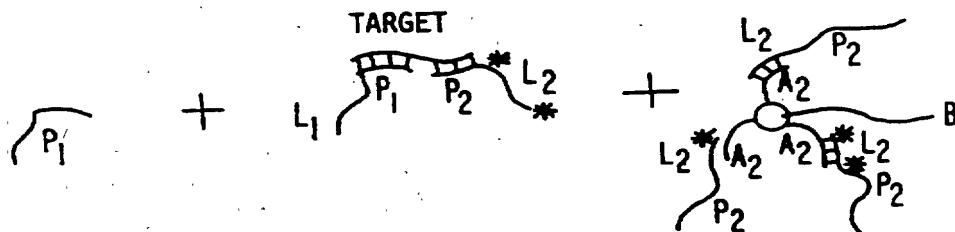
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**FIG. 3**



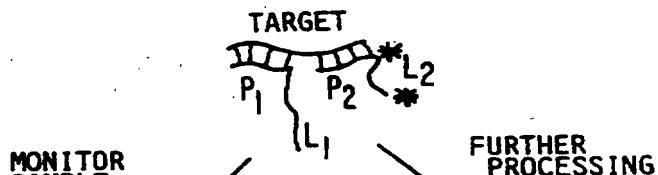
## **STEP 1**

SUPPORT SELECTIVELY BINDING  
TO SECOND LIGAND OF P<sub>2</sub>  
UNDER BINDING CONDITIONS.



## STEP 2

**REMOVE RETRIEVABLE  
SUPPORT (B<sub>1</sub>)**



**MONITOR  
SAMPLE  
FOR LABEL**

## FURTHER PROCESSING

### **STEP 3**

FIG. 4

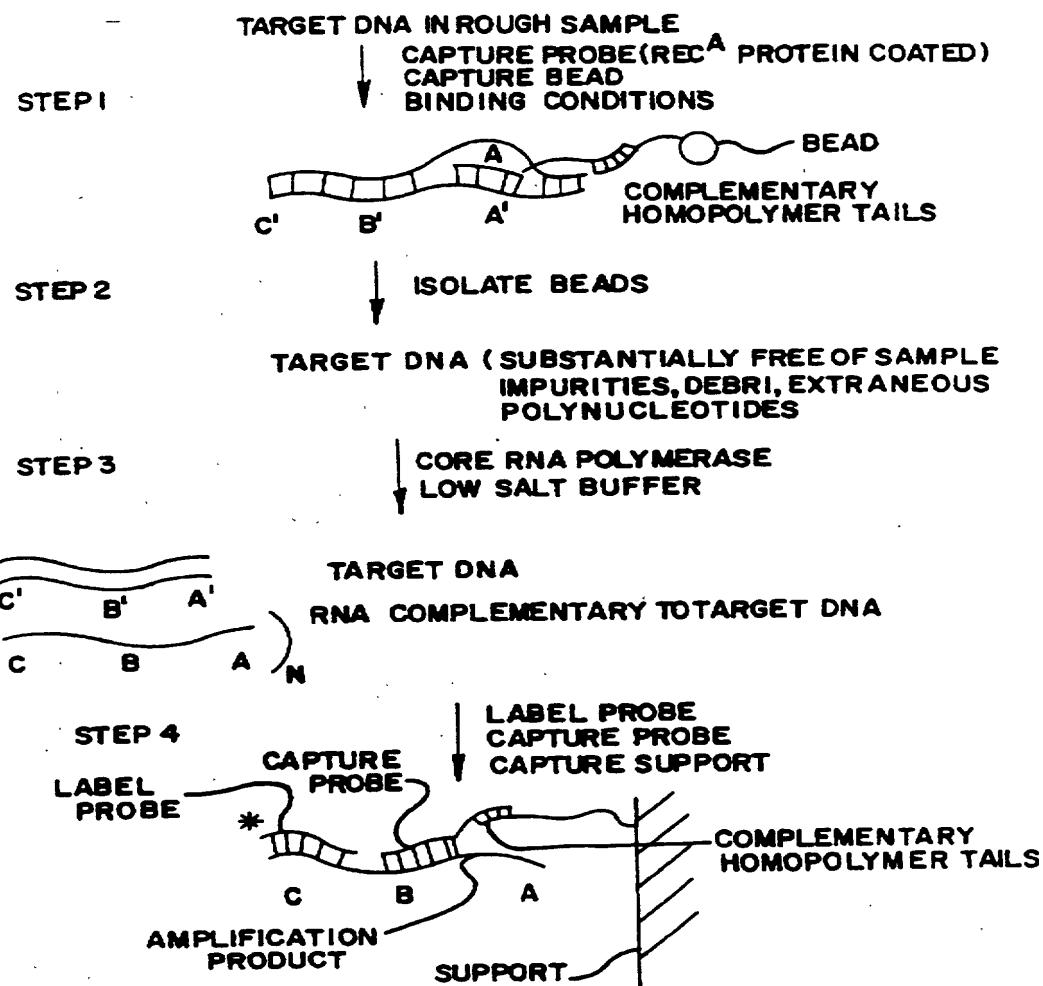
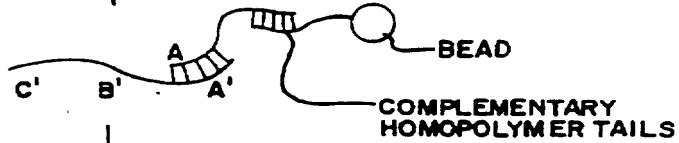


FIG. 5

TARGET DNA IN ROUGH SAMPLE  
↓  
CAPTURE PROBE  
↓  
CAPTURE BEAD

## **STEP 1**

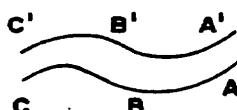


## **STEP 2**

**TARGET DNA (SUBSTANTIALLY FREE OF  
SAMPLE IMPURITIES, DEBRI,  
EXTRANEous POLYNUCLEOTIDES)**

### **STEP 3g**

## DNA POLYMERASE HEXAMER PRIMERS



STEP 3b

**CORE RNA POLYMERASE  
LOW SALT BUFFER**



## **STEP 4**

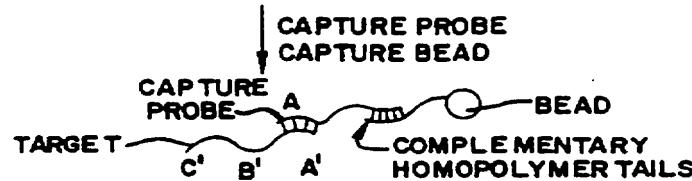
## LABELLED PROBE CAPTURE PROBE SUPPORT

The diagram illustrates a hybridization assay setup. A vertical line on the right represents the support. A capture probe (A) is attached to the support and binds to a target sequence (B). A label probe (C) is attached to the target sequence B. Its complementary sequence C' is attached to a second capture probe (A') on the support. The label probe C is labeled with an asterisk (\*).

FIG. 6

TARGET DNA IN ROUGH SAMPLE

STEP 1



STEP 2

ISOLATE BEAD

TARGET DNA (SUBSTANTIALLY FREE OF SAMPLE IMPURITIES, DEBRI, AND EXTRANEous POLYNUCLEOTIDES)

STEP 3a

DNA POLYMERASE  
HEXAMER PRIMERS



STEP 3b

I. DENATURE  
2. DNA POLYMERASE



FIRST COMPLEMENT  
COMPLEMENT TO FIRST

STEP 4

DENATURE  
LABEL PROBE  
CAPTURE SUPPORT  
CAPTURE PROBE

TARGET/AMPLIFICATION PRODUCT

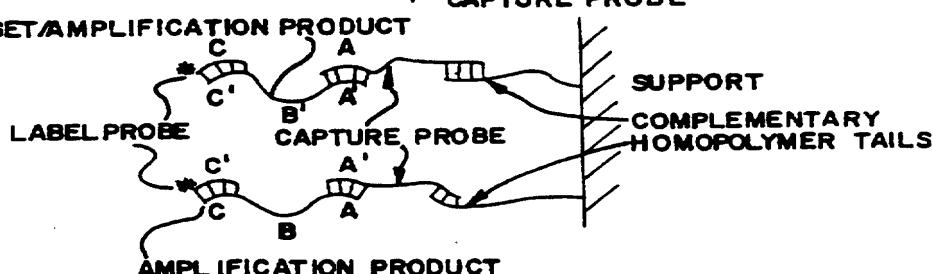
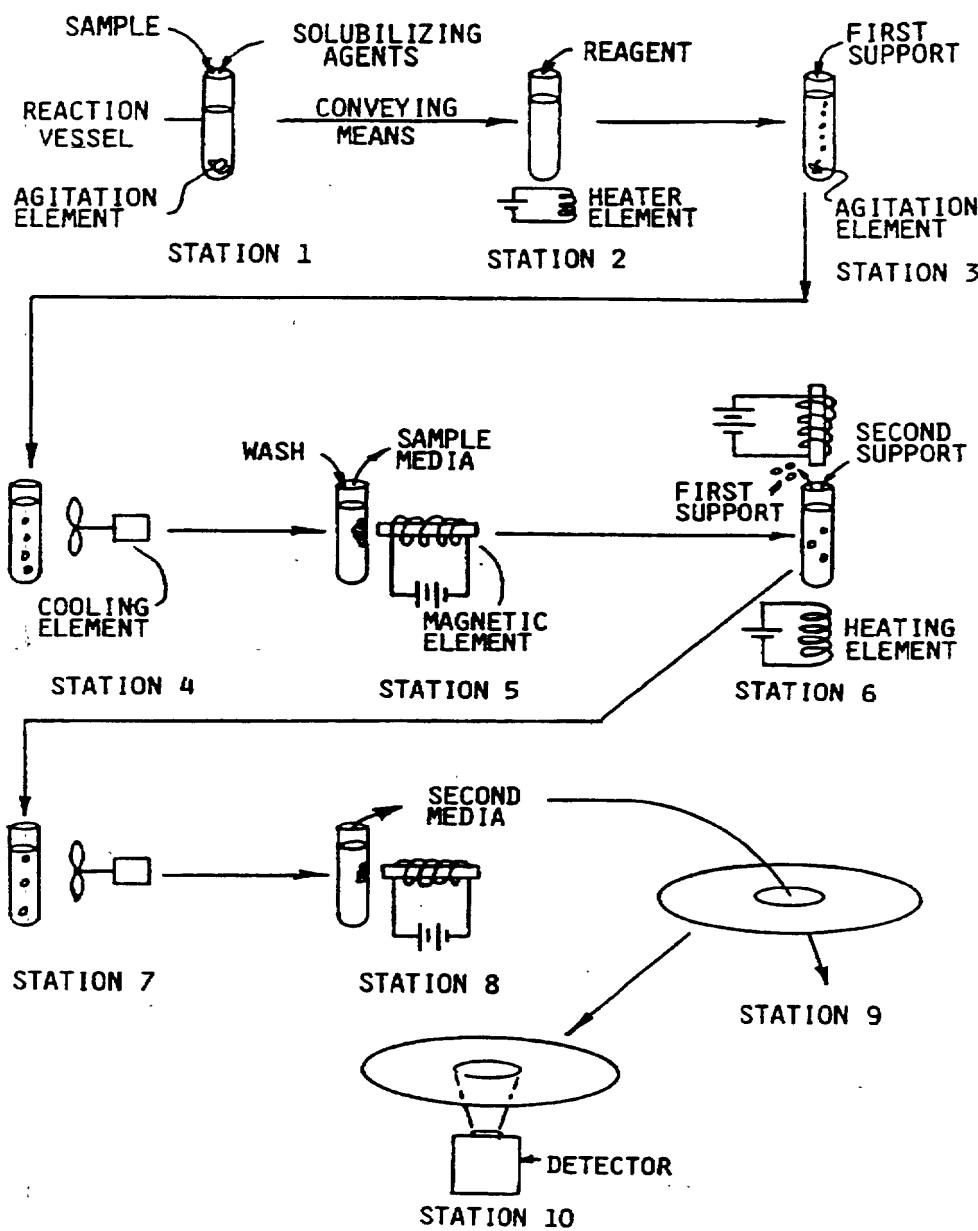
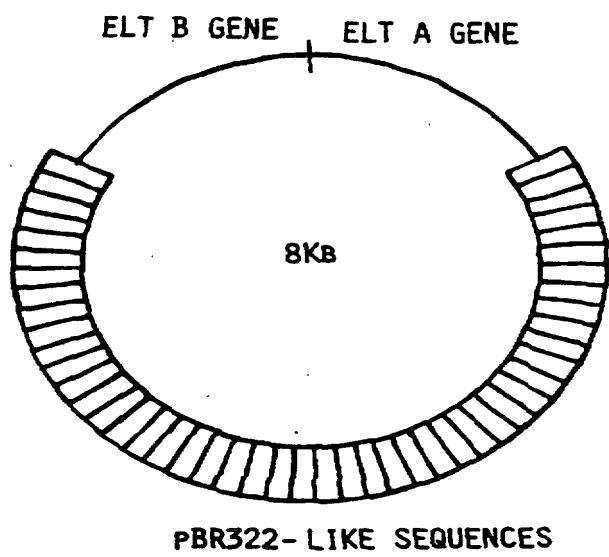


FIG. 7



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**FIG. 8**

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USA	121,935,11-US	CON	HNG	PROSECUTN	09/23/0,00	5/3/04				TACKET AND BACKGROUNDS CARRYING METHODS WITH APPLICATION FOR SECURITY ASSAY	COLUMBUS	AC

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FIG. 1a

CELLS  
WITHIN  
CLINICAL  
SAMPLE

STEP 1



SOLUBLIZING AGENTS  
& DENATURATION  
OF NUCLEIC ACID

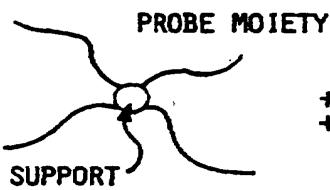
SAMPLE NUCLEIC ACID & CELLULAR DEBRI & IMPURITIES

STEP 2

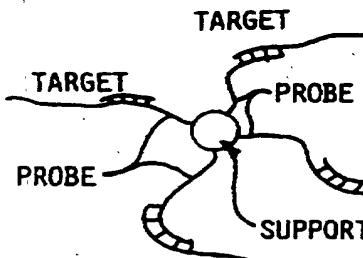
RETRIEVABLE SUPPORT  
WITH PROBE MOIETY

SAMPLE  
NUCLEIC ACID +

STEP 3



BINDING CONDITIONS

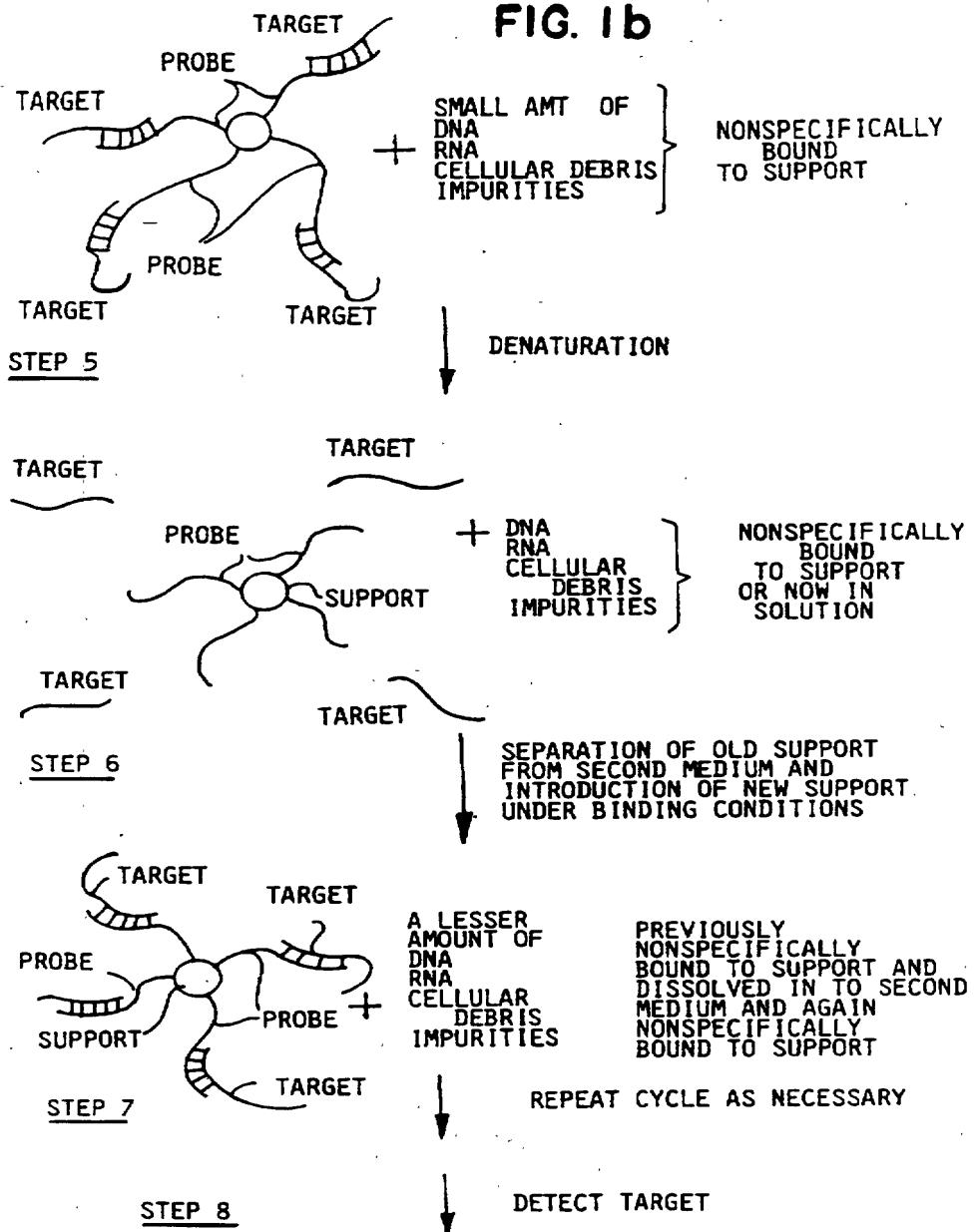


LARGE AMOUNT OF  
DNA  
RNA  
CELLULAR DEBRIS  
IMPURITIES

STEP 4

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FIG. 1b



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0008080 - 0005656560

FIG. 2a

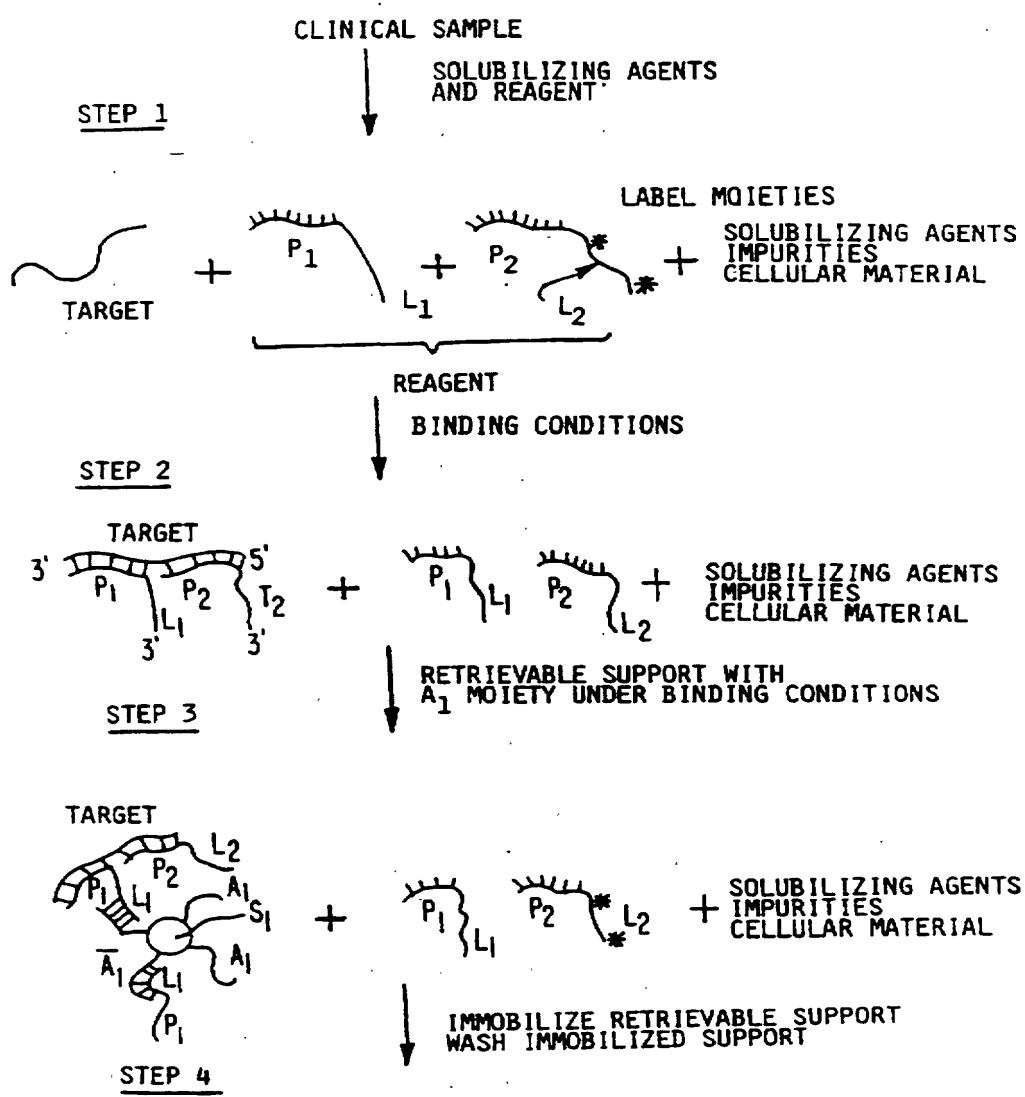


FIG. 2b

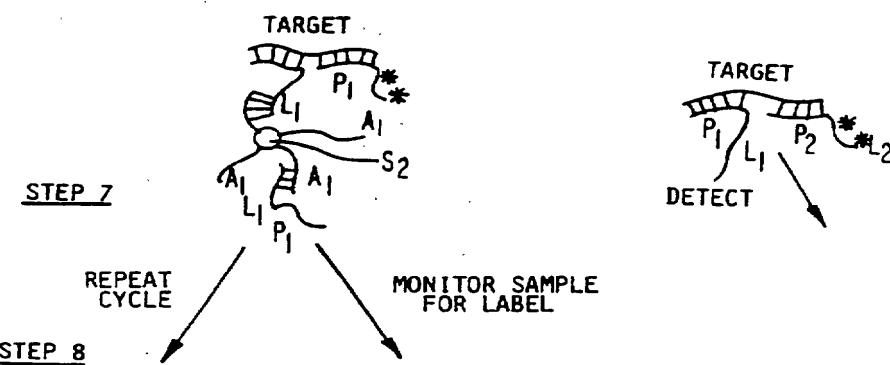
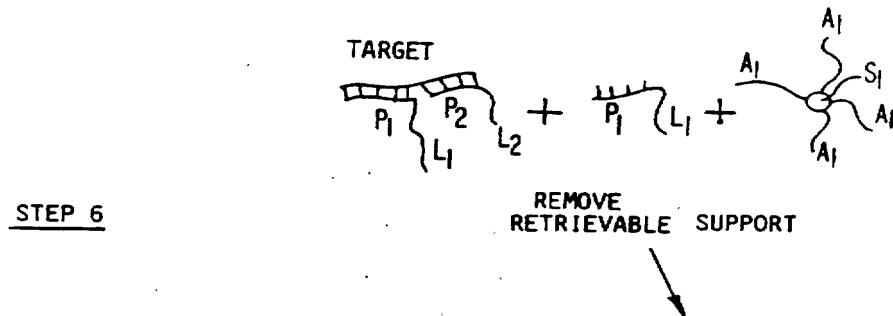
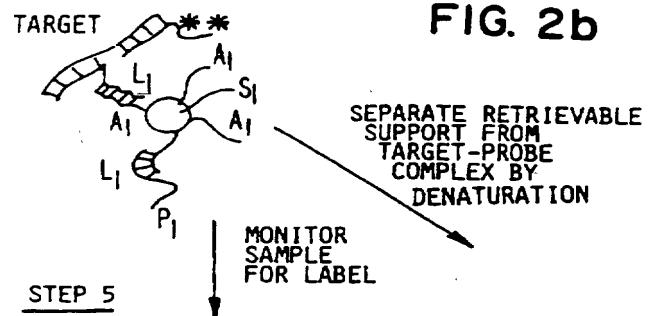
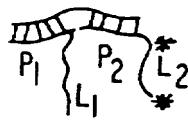
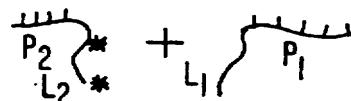


FIG. 3

TARGET



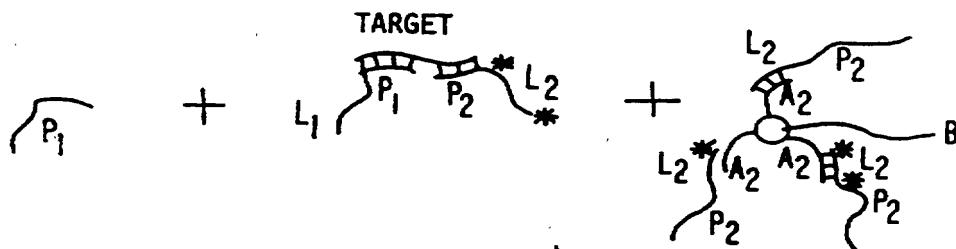
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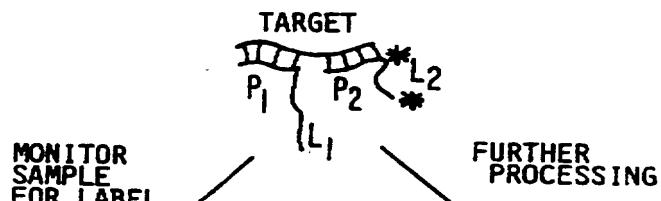
STEP 1

SUPPORT SELECTIVELY BINDING  
TO SECOND LIGAND OF  $P_2$   
UNDER BINDING CONDITIONS



STEP 2

REMOVE RETRIEVABLE  
SUPPORT ( $B_1$ )



STEP 3

MONITOR  
SAMPLE  
FOR LABEL

FURTHER  
PROCESSING

FIG. 4

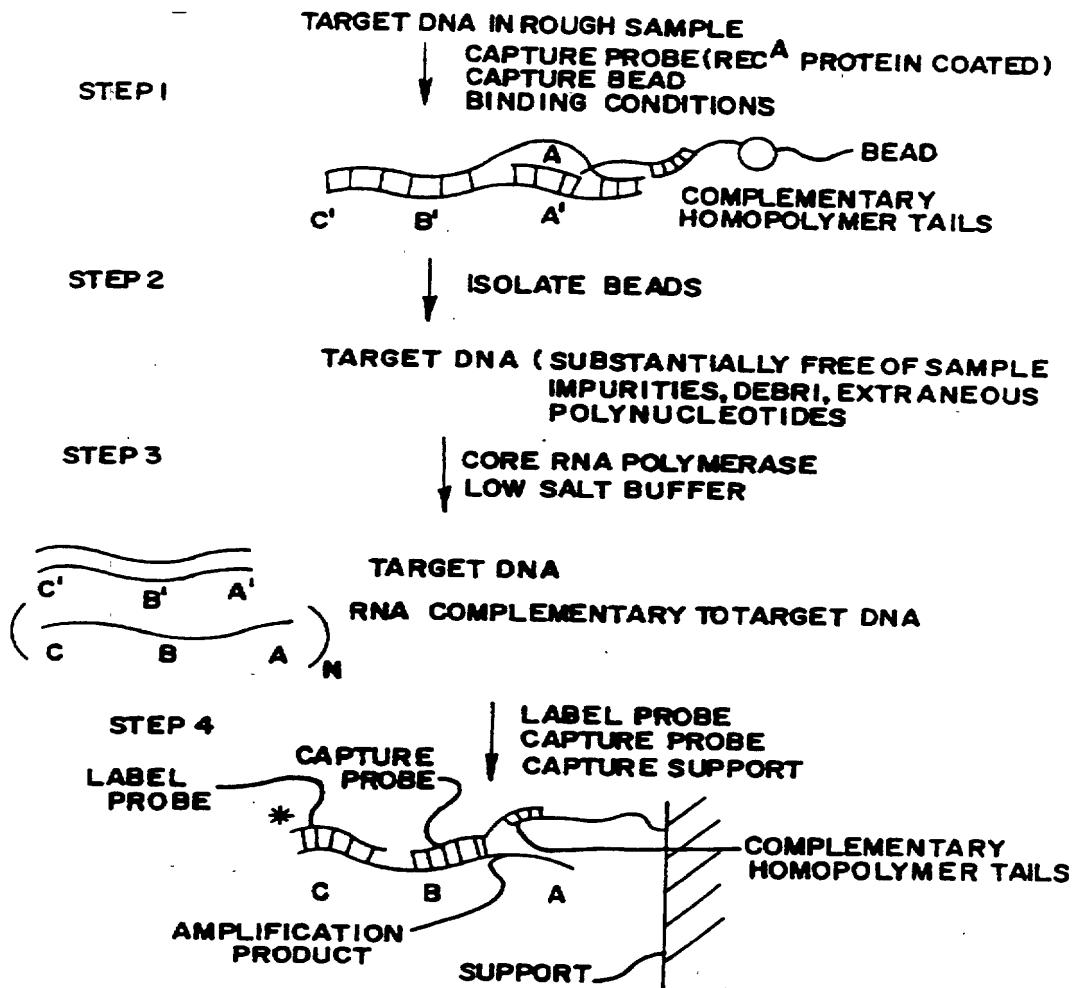


FIG. 5

TARGET DNA IN ROUGH SAMPLE  
CAPTURE PROBE  
CAPTURE BEAD

## **STEP 1**

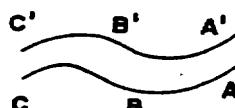


## **STEP 2**

**TARGET DNA (SUBSTANTIALLY FREE OF  
SAMPLE IMPURITIES, DEBRI,  
EXTRANEous POLYNUCLEOTIDES)**

### **STEP 3a**

## DNA POLYMERASE HEXAMER PRIMERS



### **STEP 3b**

## CORE RNA POLYMERASE LOW SALT BUFFER



#### **STEP 4**

**LABELLED PROBE  
CAPTURE PROBE  
SUPPORT**

#### **AMPLIFICATION TARGET**

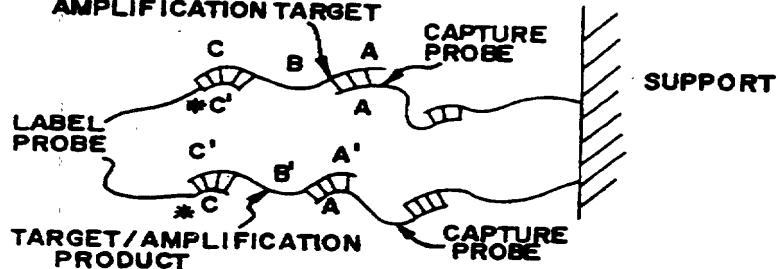
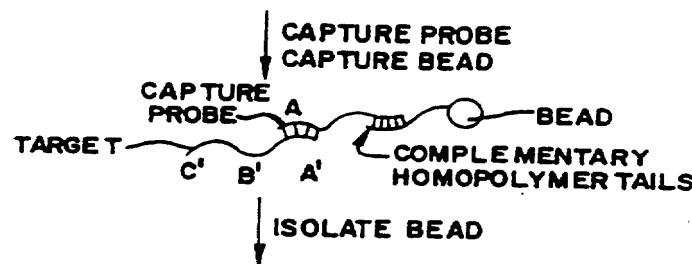


FIG. 6

TARGET DNA IN ROUGH SAMPLE

STEP 1



STEP 2

ISOLATE BEAD

TARGET DNA (SUBSTANTIALLY FREE OF SAMPLE IMPURITIES, DEBRI, AND EXTRANEOUS POLYNUCLEOTIDES)

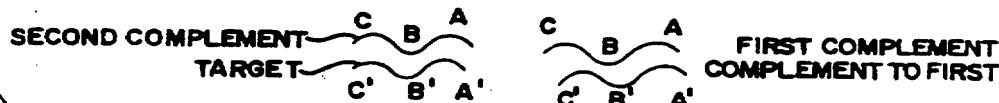
STEP 3a

DNA POLYMERASE  
HEXAMER PRIMERS



STEP 3b

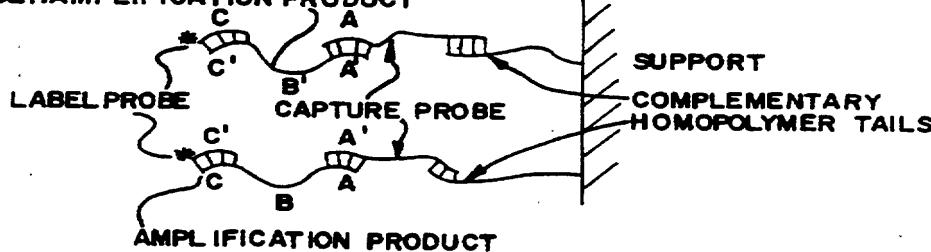
1. DENATURE  
2. DNA POLYMERASE



STEP 4

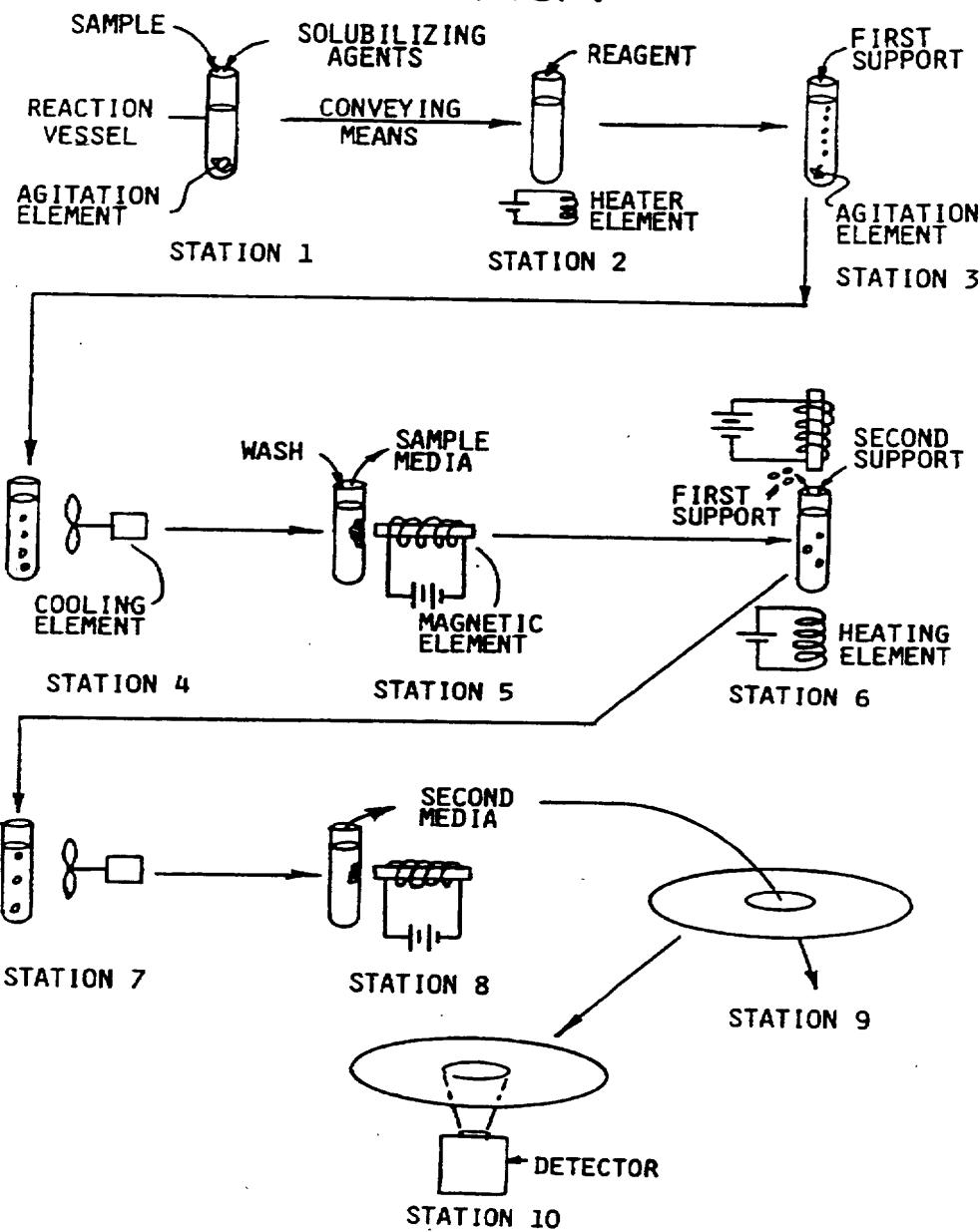
DENATURE  
LABEL PROBE  
CAPTURE SUPPORT  
CAPTURE PROBE

TARGET/AMPLIFICATION PRODUCT

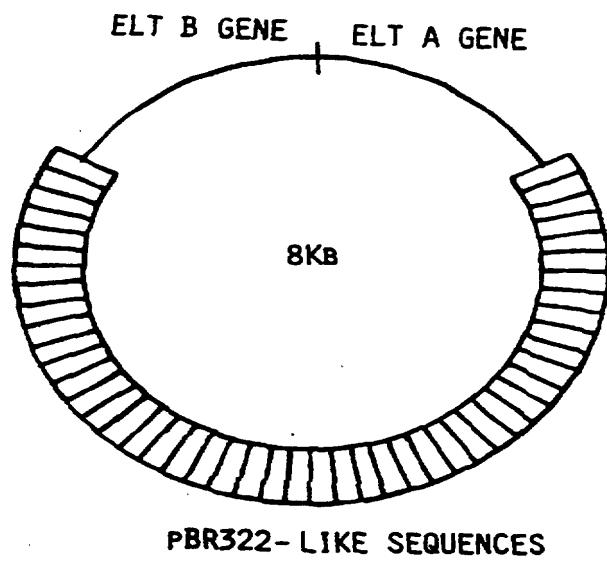


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**FIG. 7**



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**FIG. 8**

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Country	District	Type	Army.	Status	Agents	Arrival Date	Project	From Date	To Date	Notes	Link	Is it permanent	Comments
Uganda	Karamoja	Mobile	NA	Active	1	2023-01-01	Project A	2023-01-01	2023-06-30	Initial deployment	<a href="#">View</a>	Yes	Initial deployment phase for Project A.

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USA	25,835,114-3	CON HQD	PROSECUTN	00238,000	5/294					IMAGING AND BACKGROUND CAPTURE METHODS WITH AMPLIFICATION		

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**SCHEDULE 1**

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